

GRAPHICAL ABSTRACTS

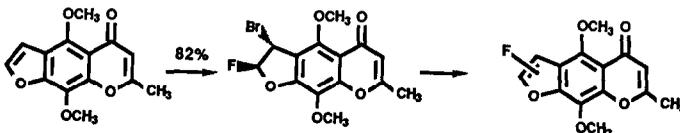
Tetrahedron Lett. 28, 4003 (1987)

THE SYNTHESIS OF 2- AND 3-FLUOROKHELLIN

S.A. Nash and R.B. Gammill

The Upjohn Company, Kalamazoo, Mich. 49001

Bromofluorination of khellin provides a fluorine containing intermediate which can be readily converted into both 2- and 3-fluorokhellin.



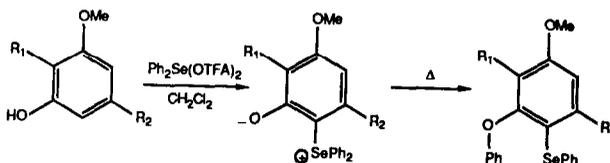
Tetrahedron Lett. 28, 4007 (1987)

SYNTHESIS AND REARRANGEMENTS OF ortho-SELENIUM PHENOXIDES

J. P. Marino* and Scott Dax

Department of Chemistry, The University of Michigan, Ann Arbor, MI 48109

A new class of selenonium phenoxides is prepared from diphenyl selenium bis(trifluoroacetate) and o-substituted phenols. These o-selenonium zwitterions undergo a thermal rearrangement to produce diaryl ethers.



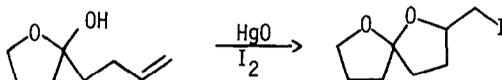
Tetrahedron Lett. 28, 4011 (1987)

ALKOXY RADICALS IN ORGANIC SYNTHESIS. A NOVEL APPROACH TO SPIROKETALS

George A. Kraus* and Jeff Thurston

Department of Chemistry, Iowa State University, Ames, Iowa 50011

Photolysis of an unsaturated hemiketal with HgO and I₂ affords a spiroketal.

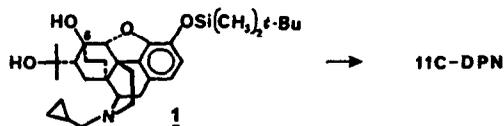


Tetrahedron Lett. 28, 4015 (1987)

SYNTHESIS OF CARBON-11 LABELED DIPRENORPHINE: A RADIOLIGAND FOR POSITRON EMISSION TOMOGRAPHIC STUDIES OF OPIATE RECEPTORS

John R. Lever,* Robert F. Dannals, Alan A. Wilson, Hayden T. Ravert and Henry N. Wagner, Jr. Departments of Environmental Health Sciences and Radiology, The Johns Hopkins Medical Institutions, Baltimore, MD 21205, USA.

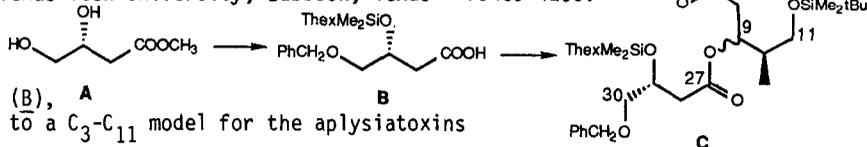
Selective alkylation of **1** with [¹¹C]-MeI followed by desilylation affords [¹¹C]-diprenorphine ([¹¹C]-DPN) labeled at the 6-methoxy position in 10% yield with a specific activity of 1740 mCi/μmol at the end of a 30 minute synthesis.



STUDIES ON THE SYNTHESIS OF THE APLYSIATOXINS: SYNTHESIS OF A SELECTIVELY-PROTECTED FORM OF THE C₂₇-C₃₀ (DIHYDROXY-BUTANOATE) MOIETY OF OSCILLATOXIN A

Robert D. Walkup* and Raymond T. Cunningham - Department of Chemistry & Biochemistry, Texas Tech University, Lubbock, Texas 79409-4260.

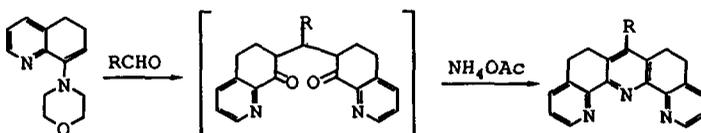
Selective benzylation of diol A via BnBr/Ag₂O/ultrasound led to the title compound (B), A which cleanly coupled to a C₃-C₁₁ model for the aplysiatoxins to form the ester C.



POLYAZA CAVITY SHAPED MOLECULES 13. A VERSATILE NEW TERPYRIDINE SYNTHESIS

Vidyadhar Hegde, Yurrgdong Jehng, and Randolph P. Thummel*
Department of Chemistry, University of Houston, Houston, Texas 77004

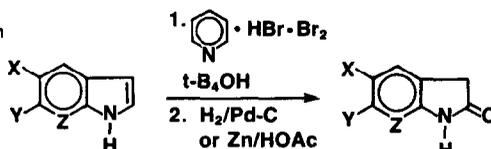
The reaction of morpholine enamines derived from α -pyridyl ketones with aromatic aldehydes leads to the formation of a 1,5-diketone intermediate which in turn provides a 4'-substituted derivative of 2,2':6'2"-terpyridine.



OXIDATION OF INDOLES WITH PYRIDINIUM BROMIDE PERBROMIDE A SIMPLE AND EFFICIENT SYNTHESIS OF 7-AZAOXINDOLES

Anthony Marfat* and Michael P. Carta
Central Research, Pfizer Inc., Groton, CT 06340

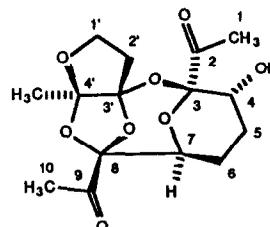
Indoles and azaindoles on treatment with pyridinium bromide perbromide in t-butanol give 3,3-dibromooxindoles which, upon further hydrogenation or reduction, yield the corresponding oxindoles.



A NOVEL TETRACYCLIC POLYKETAL FROM THE MARINE RED ALGA LAURENCIA CHILENSIS

M. Bittner, F. Gonzalez, H. Valdebenito, and M. Silva*, Laboratorio de Quimica de Productos Naturales, Universidad de Concepcion, Casilla 2407, Apartado 10, Concepcion, CHILE
V.J. Paul and W. Fenical*, Scripps Institution of Oceanography, La Jolla, CA 92093
M.H.M. Chen and Jon Clardy*, Department of Chemistry-Baker Laboratory, Cornell University, Ithaca, NY 14853-1301

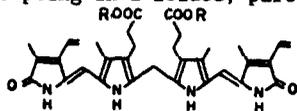
An unusual tetracyclic polyketal (1) was isolated from the red alga *Laurencia chilensis*. Structure 1 was established by x-ray diffraction and spectral methods.



CONFORMATION OF BILIRUBIN ESTERS FROM CD SPECTROSCOPY

David A. Lightner*, Man-Hua Zhang and Francesc R. Trull
 Chemistry Department, University of Nevada, Reno, NV 89557

Optically active (4*Z*,15*Z*)-bilirubin-IX α mono and diesters were prepared from the methanesulfonate of β -(+)-2-butanol. The more intense bisignate CD of the mono-ester is characteristic of exciton coupling in a folded, partially intramolecularly H-bonded conformation.



monoester: $\Delta\epsilon_{\text{max}}^{405} +10.2$, $\Delta\epsilon_{\text{max}}^{450} -15.7$ (CH_2Cl_2)

diester: $\Delta\epsilon_{\text{max}}^{400} +1.3$, $\Delta\epsilon_{\text{max}}^{450} -1.7$ (CH_2Cl_2)

DIASTERESELECTIVITY IN THE DIRECTED ALDOL
 CONDENSATION OF 2-TRIMETHYLSILOXYFURAN WITH

ALDEHYDES. A STEREODIVERGENT ROUTE TO THREO AND ERYTHRO δ -HYDROXY- γ -LACTONES

Charles W. Jefford, Danielle Jaggi, and John Boukouvalas,
 Department of Organic Chemistry, University of Geneva, 1211 Geneva 4, Switzerland

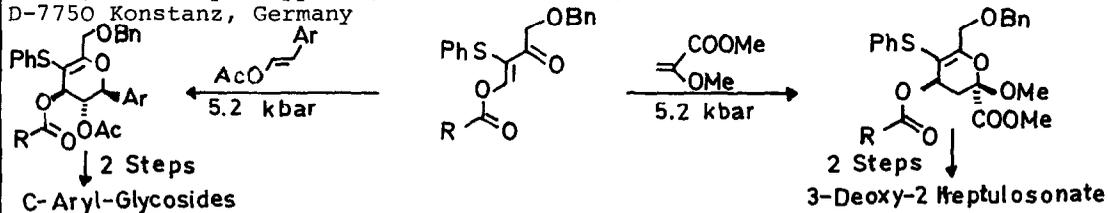
THE SYNTHESIS OF (\pm)-CAVERNOSINE

Charles W. Jefford, Danielle Jaggi, Gérald Bernardinelli and John Boukouvalas,
 Department of Organic Chemistry and Laboratory of Crystallography, University of
 Geneva, 1211 Geneva 4, Switzerland



C-ARYL-GLYCOSIDES AND 3-DEOXY-2-GLYCULOSONATES
 VIA INVERSE TYPE HETERO-DIELS-ALDER REACTION

Richard R. Schmidt*, Wendelin Frick, Brigitte Haag-
 Zeino, and Satyam Apparao, Fakultät Chemie, Universität Konstanz, Postfach 5560,
 D-7750 Konstanz, Germany

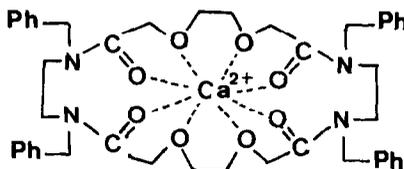


**UN TETRALACTAME MACROCYCLIQUE DERIVE DE
L'ACIDE DIOXA-3.6 OCTANEDIOIQUE,
NOUVEL IONOPHORE NEUTRE SELECTIF DU CALCIUM**

Nadine LEYGUE, Louis CAZAUX, Claude PICARD and Pierre TISNES

Synthèse et Physicochimie organique, UA CNRS N° 471,
Université Paul Sabatier - 118, route de Narbonne
31062 TOULOUSE CEDEX (FRANCE)

Synthetic procedure to the title compound
and its complexation ability and selectivity
forward Ca^{2+} is described

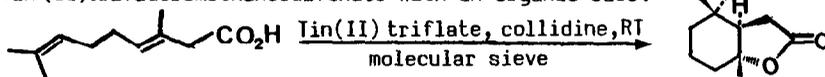


**CYCLISATION ELECTROPHILE INDUITE PAR LE TRIFLATE STANNEUX :
SYNTHESE DU (±)-TETRAHYDROACTINIDIOLIDE TRANS**

Noukpo GNONLONFOUN et Henri ZAMARLIK

Laboratoire de Chimie Organique et Biologique - Faculté des Sciences et Techniques
Université de TOURS Parc de Grandmont 37200 TOURS FRANCE

Title compound was prepared by a one pot reaction starting from homogeric acid and using
Tin(II)trifluoromethanesulfonate with an organic base.



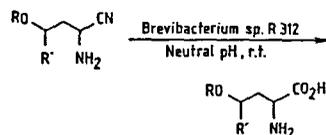
**BACTERIA IN ORGANIC SYNTHESIS : γ -ALKOXY- α -AMINOACIDS FROM RELATED
 α -AMINONITRILES.**

Yen Vo-Quang^{*1}, Dominique Marais¹, Liliane Vo-Quang¹ and François Le Goffic¹.

Alain Thiéry², Marc Maestracci², Alain Arnaud² and Pierre Galzy².

¹ Laboratoire de Bioorganique et Biotechnologies, Ecole Nationale Supérieure
de Chimie de Paris, 11 rue P. et M. Curie 75231 PARIS Cédex 05 (France).

² Chaire de Génétique et Microbiologie, Ecole Nationale Supérieure Agronomique,
place P. Viala 34060 MONTPELLIER Cedex (France).

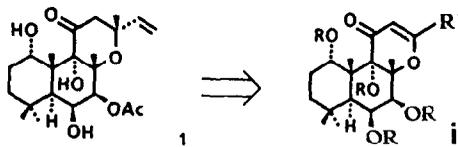


RETROSYNTHETIC STUDIES WITH FORSKOLIN

Bernard Delpech and Robert Lett

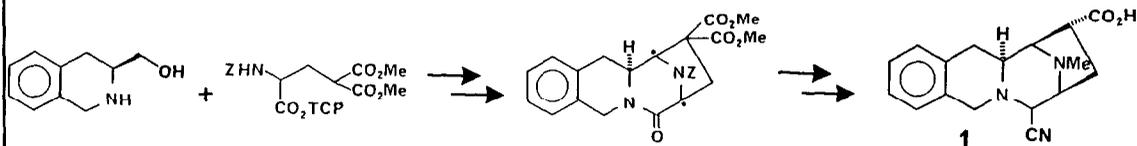
Laboratoire CNRS/ROUSSEL UCLAF - BP 9 - 93230 Romainville (France)

New preparations of some dihydro- γ -pyrones **1** derived from
forskolin **1** are reported. Conjugate addition of cuprates is
shown to occur on **i**, in good yield, in the presence of
 $\text{BF}_3 \cdot \text{Et}_2\text{O}$: dimethyl and vinyl cuprates add with major α
stereoselectivity, whereas dibutyl cuprate adds mostly β .



SYNTHETIC APPROACH TO QUINOCARCIN

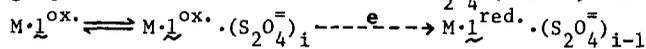
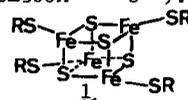
Hiromitsu Saito and Tadashi Hirata*

Tokyo Research Laboratories, Kyowa Hakko Kogyo Co., Ltd.
3-6-6, Asahi-machi, Machida-shi, Tokyo, Japan

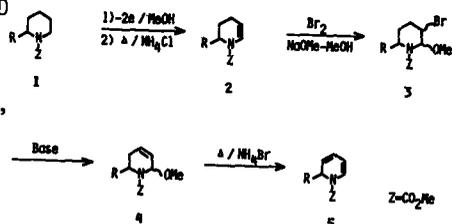
A REVERSIBLE AND SPECIFIC MULTIPLE BINDING OR ADSORPTION OF DITHIONITE ANION ON CATIONIC MICELLAR SURFACE.

Iwao Tabushi, Yasuhisa Kuroda*, Yoro Sasaki

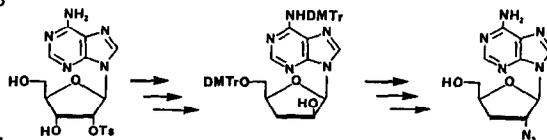
Department of Synthetic Chemistry, Kyoto University, Sakyo-ku, Kyoto, 606, Japan

The kinetic analysis of the reduction of 1 in aq. CTAB- $S_2O_4^{2-}$ solution indicated the multiple binding equilibrium between micelles and $S_2O_4^{2-}$ ($n=19\pm 7, K=1300\pm 400M^{-1}, k=2200\pm 300M^{-1/2}\cdot s^{-1}$).M; CTAB micelle, $i=1-n$. $R=CH_2CH(CO_2Na)NHCOC_{17}H_{35}-n-L$ 

A NEW METHOD FOR REGIOSELECTIVE SYNTHESIS OF 2-SUBSTITUTED 1-(METHOXYCARBONYL)-1,2-DIHYDROPYRIDINES

Tatsuya Shono,* Yoshihiro Matsumura, Osamu Onomura,
and Yasufu YamadaDepartment of Synthetic Chemistry, Faculty of Engineering,
Kyoto University, Yoshida, Sakyo, Kyoto 606, JAPANA facile synthesis of 2-substituted 1-(methoxycarbonyl)-
1,2-dihydropyridines 5 from 2-substituted piperidines 1.

THE FACILE SYNTHESIS OF 2'-AZIDO-2',3'-DIDEOXYADENOSINE.

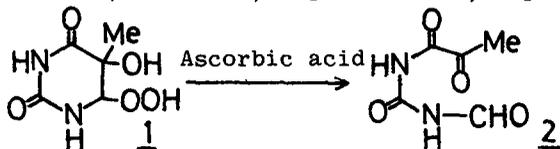
PREPARATIVE APPLICATIONS OF THE DEOXYGENATIVE
[1,2]-HYDRIDE SHIFT AND β -ELIMINATION REACTIONS
OF SULFONATES WITH $Mg(OMe)_2-NaBH_4$ Masajiro Kawana* and Hiroyoshi Kuzuhara
RIKEN (The Institute of Physical and Chemical
Research), Wako-shi, Saitama 351-01, JapanThe title compound was prepared from 2'- or 3'-
O-tosyladenosine in 37 or 51% overall yield, respectively.

AN UNUSUAL REACTION OF CIS-6-HYDROPEROXY-5-HYDROXY-5,6-DIHYDROTHYMINE WITH ASCORBIC ACID

Toshio Itahara

College of Liberal Arts, Kagoshima University, Korimoto, Kagoshima 890, Japan

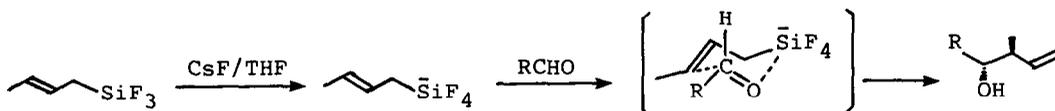
CIDNP was observed on reaction of 1 with ascorbic acid in D₂O at room temperature to give 2.



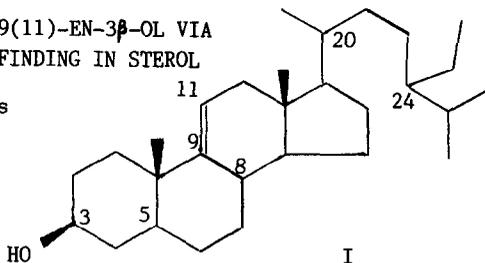
REGIOSPECIFIC AND HIGHLY STEREOSELECTIVE ALLYLATION OF ALDEHYDES WITH ALLYLTRIFLUOROSILANE ACTIVATED BY FLUORIDE IONS

Mitsuo Kira,* Mineo Kobayashi, and Hideki Sakurai*

Department of Chemistry, Tohoku University, Sendai 980, Japan

DIRECT CYCLISATION OF SQUALENE TO 5 α -STIGMAST-9(11)-EN-3 β -OL VIA $\Delta^{9(11)}$ LANOSTEROL IN *COSTUS SPECIOSUS*: A UNIQUE FINDING IN STEROL BIOSYNTHESIS. A. AKHILA, M.M. GUPTA & R.S. THAKURCentral Institute of Medicinal & Aromatic Plants
Lucknow, India 226 016

Squalene metabolises into 5 α -stigmast-9(11)-en-3 β -ol (I) via $\Delta^{9(11)}$ lanosterol rather than $\Delta^{8(9)}$ lanosterol. A report for the first time in sterol biosynthesis in higher plants.

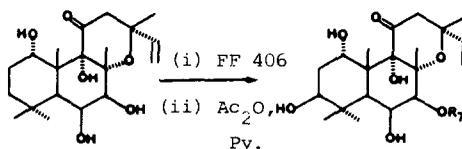


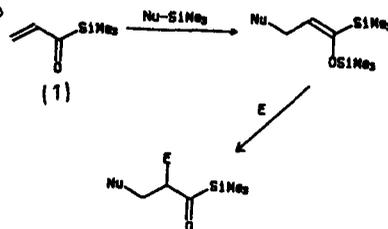
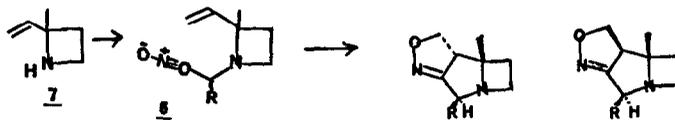
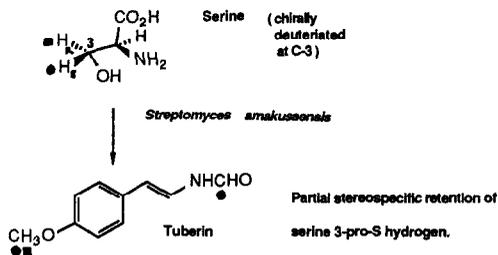
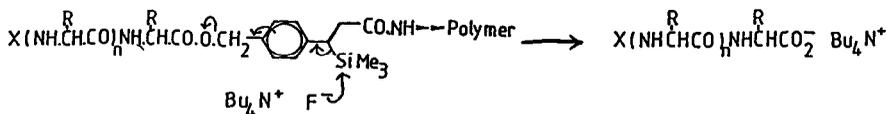
SYNTHESIS OF METABOLITES OF FORSKOLIN!

Y. Khandelwal*, N.J. de Souza, S. Chatterjee, B.N. Ganguli
and R. H. Rupp

Centre For Basic Research, Hoechst India Limited, Bombay 400 080, INDIA.

Microbial and chemical transformations of 7-deacetylforskolin are described leading to 3 β -hydroxy forskolin and its derivatives which have been found to be metabolites of forskolin.



Tetrahedron Lett. 28,4093 (1987)SYNTHESIS OF POLYFUNCTIONALIZED ACYLSILANES
VIA PROPENOYLTRIMETHYLSILANEA. Ricci, A. Degl'Innocenti, G. Borselli and G. Reginato
Centro CNR Composti Eterociclici, c/o Dipartimento
di Chimica Organica, via Capponi 9, Firenze, Italy.Propenoyltrimethylsilane (1) reacts with a wide series of silylated nucleophiles affording β -functionalized silyl enol ethers of acylsilanes: "in situ" reaction of these compounds leads to polyfunctionalized acylsilanes.Tetrahedron Lett. 28,4097 (1987)STEREOSELECTIVITY DURING NITRILE OXIDE-OLEFIN CYCLOADDITIONS.
SYNTHESIS OF FUSED AZETIDINESAlfred Hassner* and K.S. Keshava Murthy
Department of Chemistry, Bar-Ilan University, Ramat-Gan 52100, IsraelTwo routes are described for converting vinyl azetidine 7 into the nitrile oxides 6, which undergo stereoselective dipolar cycloaddition to fused azetidines.Tetrahedron Lett. 28,4101 (1987)STEREOCHEMISTRY ASSOCIATED WITH
THE CONVERSION OF SERINE THROUGH
METHYLENETETRAHYDROFOLATE AND
METHENYLTETRAHYDROFOLATE IN THE
BIOSYNTHESIS OF TUBERINK.M. Cable & R.B. Herbert, Dept. of Organic
Chemistry, The University, Leeds LS2 9JT
V. Bertram & D.W. Young, School of Chemistry
and Molecular Sciences, University of Sussex,
Falmer, Brighton BN1 9QJ.SOLID PHASE PEPTIDE SYNTHESIS:
FLUORIDE ION RELEASE FROM THE RESIN.R. RAMAGE*, C.A. BARRON, S. BIELECKI and D.W. THOMAS
Department of Chemistry, University of Edinburgh,
West Mains Road, Edinburgh, EH9 3JJ, Scotland, UKSynthesis of a linker for solid phase peptide synthesis which allows release by $Bu_4N^+F^-$ of the product peptideTetrahedron Lett. 28,4105 (1987)

SYNTHESIS, STRUCTURE AND REACTIVITY OF 4-ACETYLAMINO-9a-ACETOXY-1,9,10-ANTHRACENETRIONE

Francisco Farfía, M. Carmen Paredes and Valter Stefani

*Instituto de Química Orgánica General, C.S.I.C., 28006 Madrid, Spain, and**Instituto de Química U.F.R.G.S., 90.000 Porto Alegre (RS), Brasil*

The synthesis, rearrangement, cycloaddition of cyclopentadiene and reaction with nucleophiles of the title compound are reported.

